EU Declaration of Conformity

Manufacturer's Name:	Van der Bend B.V.
Manufacturer's Address:	Kloosterweg 34, 3232 LC Brielle (NL)
SRN (Single Registration Number):	NL-MF-000001575
Basic UDI-DI:	871705672460089
Name of the Device(s):	Van der Bend Patch Test Chambers
Classification:	Class I

Conformity assessment route:

The Van der Bend Patch Test Chambers have been classified as Class I according to Annex VIII, Rule 1, and are in conformity with the general safety and performance requirements and provisions of Regulation (EU) 2017/745 on medical devices.

The device is in conformity with the relevant standards listed in the current technical documentation, including:

ISO 14971:2019
ISO 15223-1:2014
ISO 10993-5:2009
ISO 10993-10:2010

The technical documentation is maintained in accordance with Annexes II and III of Regulation (EU) 2017/745.

This EU Declaration of Conformity is issued under the sole responsibility of Van der Bend B.V.

We hereby declare that the medical device specified above meets the applicable provisions of Regulation (EU) 2017/745 on medical devices.

All supporting documentation is retained at the premises of the manufacturer.

Product reference list

Manufacturer	Product name	Article reference	Basic UDI-DI	UDI-DI (GTIN)	Classification
Van der Bend B.V.	Van der Bend Patch Test Chambers	4600	871705672460089	08717056720104	Class I (Rule 1)

Signature:

Annemarie van der Bend
Director



Place and date of issue:
Brielle, 14 September 2026